

Electro-Optical Effects in Liquid Crystals

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- I. General picture
- II. Basic effects (nematics, cholesterics, smectics)
- III. Frederiks transition: past and present
- IV. Helix unwinding: pro and contra
- V. Electro-optics of a spatial soliton: glance in future

For more details and earlier achievements see

L.M. Blinov, V.G. Chigrinov, **Electrooptic Effects in Liquid Crystal Materials**, Springer, 1996, p.464, figs. 221.

I. General picture

1. Why liquid crystals?

Gases, liquids and isotropic solids (e.g. polymers)

a) Electric field induced shift and splitting of spectral lines (Stark, 1913)

$$\Delta W = \Delta p E + \Delta \alpha E^2$$

and correspondent change in absorption and refraction (Kramers-Kronig)

b) Electric field induced reorientation of anisotropic molecules (Kerr, 1875)

In both cases: **quadratic-in-field EOE**: $\Delta n \propto k E^2$

Solid systems with polar symmetry such as polar crystals (Pockels, 1984), ferroelectrics (Valašek, 1921), or poled polymers called electrets

In this case: **linear-in-field EOE** $\Delta n \propto \beta E$

The key parameter is $\Delta n(E)$

Liquid crystals: what is new?

Large Δn ? No, some molecular crystals have Δn even larger, up to 0.5.

Low viscosity? No, many polar liquids have smaller viscosity (e.g., nitrobenzene)

New effect? No, the field induced molecular reorientation like in the Kerr effect

Really new is

$$Q = S(n_i n_j - 1/3)$$

the new type of molecular ordering in a liquid state.

Now the electric field can **reorient the director** (optical axis, i.e. “direction of Δn ”)

In the bulk, the reorientation of all molecules at once does not require energy (Goldstone mode). In a thin cell, the reorientation of almost all molecules at once requires only low energy (due to small Frank elastic moduli in the liquid state)

The EOE is strong even **in thin cells at low voltages** $I = I_0 \sin^2 \beta \sin^2 \frac{2\pi d \langle \Delta n \rangle}{\lambda}$

The switching times **in thin cells** are sufficiently short (ms) $\tau \approx \frac{\gamma d^2}{K \pi^2}$ or $\approx \frac{\gamma}{PE}$

LCs are materials for thin film devices with many pixels, like displays, SLM, etc

2. A principle of free energy minimum (with bulk + surface terms)

$$F = F_V + F_S = \int_V (g_{elast} + g_{field}) dV + F_S$$

Density of elastic energy with splay, twist and bend terms (Frank):

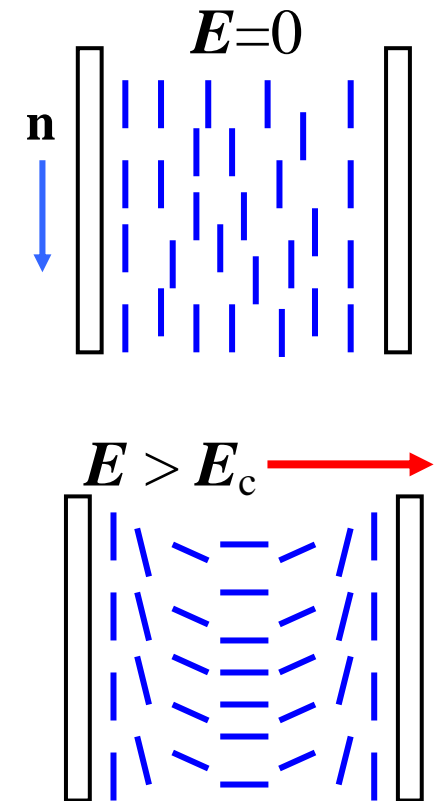
$$g_{elast} = K_1 (\text{div} \mathbf{n})^2 + K_2 (\mathbf{n} \text{curl} \mathbf{n} + q_0)^2 + K_3 (\mathbf{n} \times \text{curl} \mathbf{n})^2$$

Density of electric energy (quadratic and linear in field):

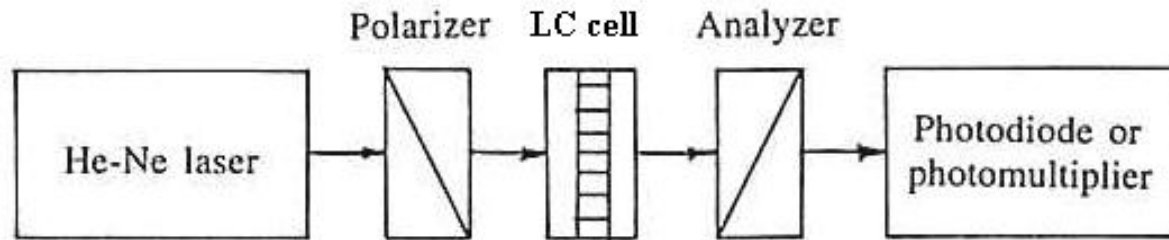
$$g_{field}^Q = -\frac{\mathbf{D} \cdot \mathbf{E}}{8\pi} = -\frac{\epsilon_{\perp} E^2}{8\pi} - \frac{\epsilon_a (\mathbf{E} \cdot \mathbf{n})^2}{8\pi}$$

$$g_{field}^L = -\mathbf{P} \cdot \mathbf{E}, \quad \mathbf{P} = \mathbf{P}_{spont} \quad \text{or} \quad \mathbf{P}_{flex}$$

$$\mathbf{P}_{flex} = e_{splay} \mathbf{n} \cdot \text{div} \mathbf{n} + e_{bend} (\mathbf{n} \times \text{curl} \mathbf{n})$$



3. Electro-optical effects (general)



Planar structures:

- (i) Transmission or reflection modes,
- (ii) any sources, white or monochromatic,
- (ii) any field, longitudinal or in-plane with interdigitated electrodes

For waveguiding geometry: come to the next lecture!

In any case we have the same list of effects:

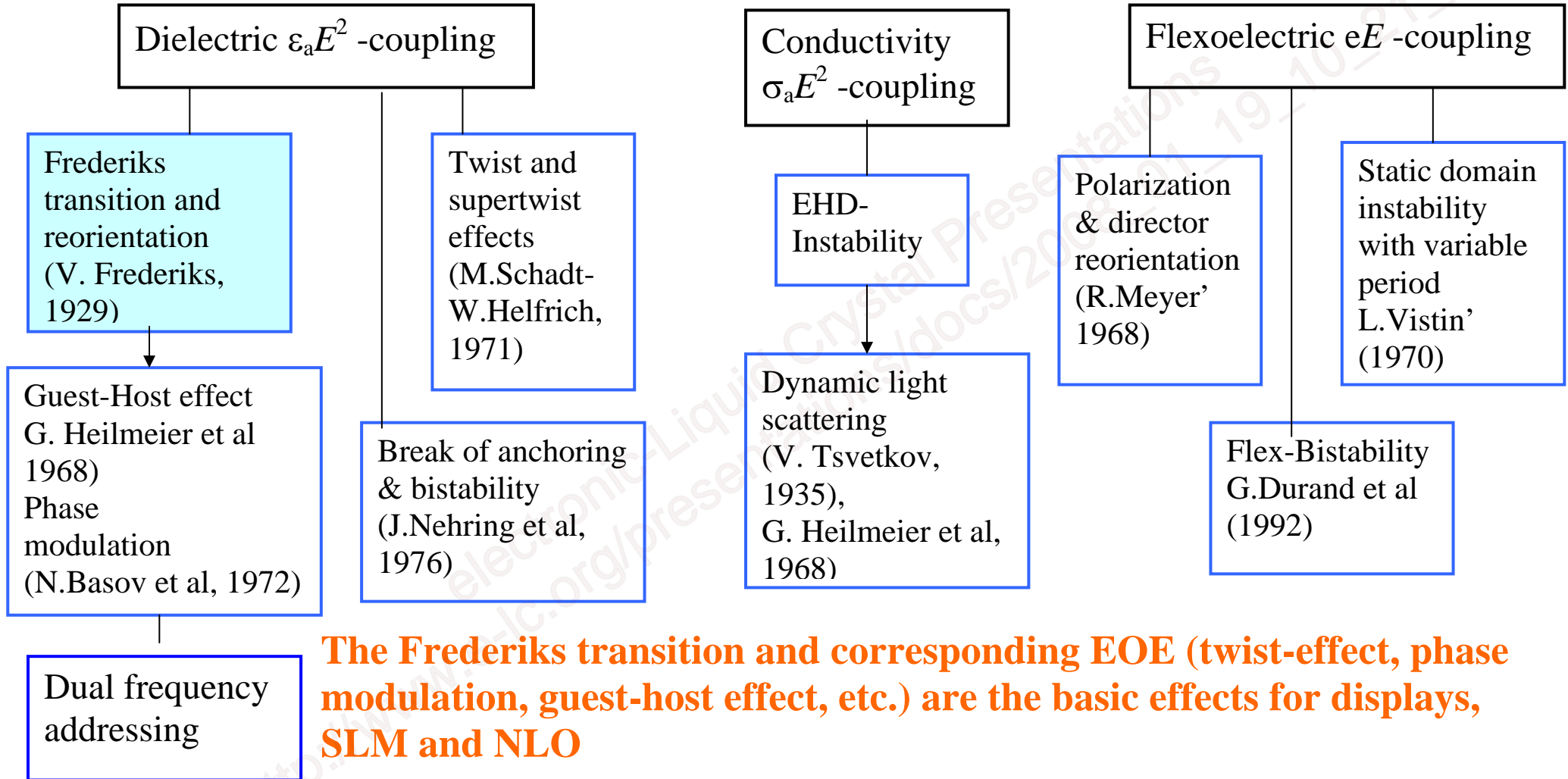
Field controlled **birefringence**

Field controlled **absorption** positive (color) or negative (lasing)

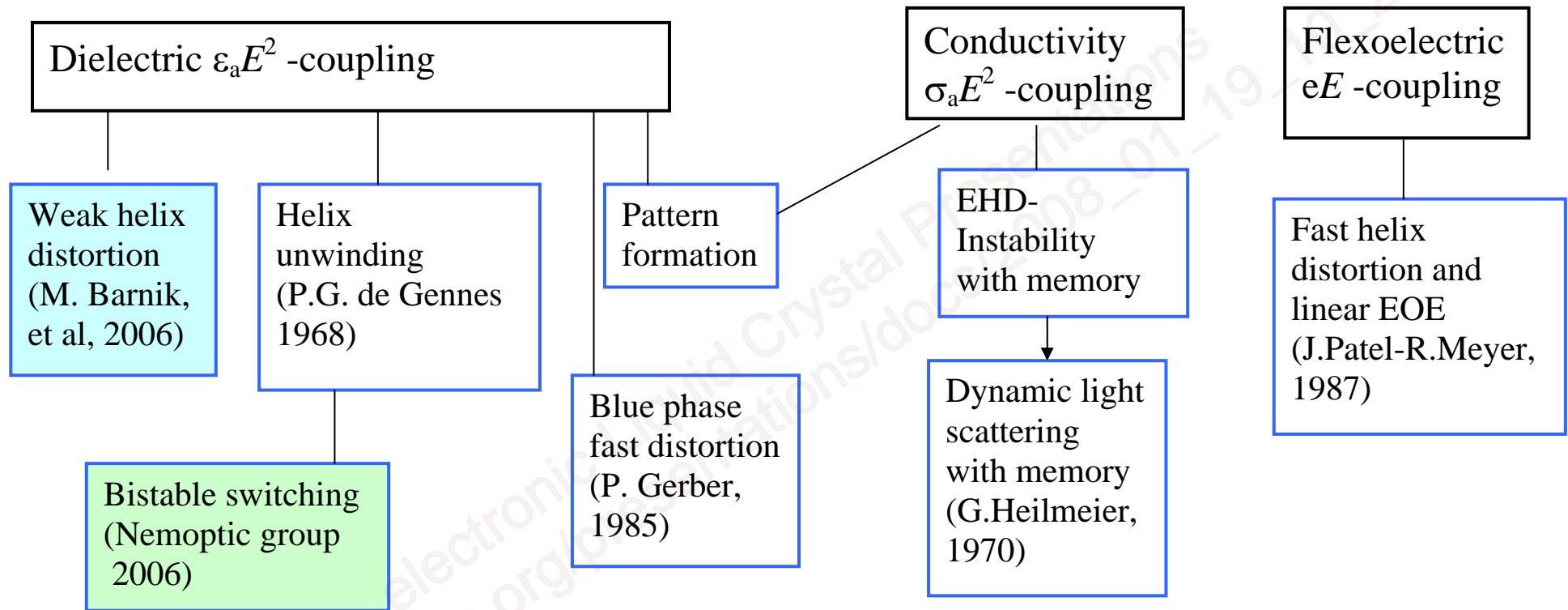
Field controlled **scattering or diffraction**

II.1 Basic EOE: Nematics

First observations: I. Bjornstahl (1918)

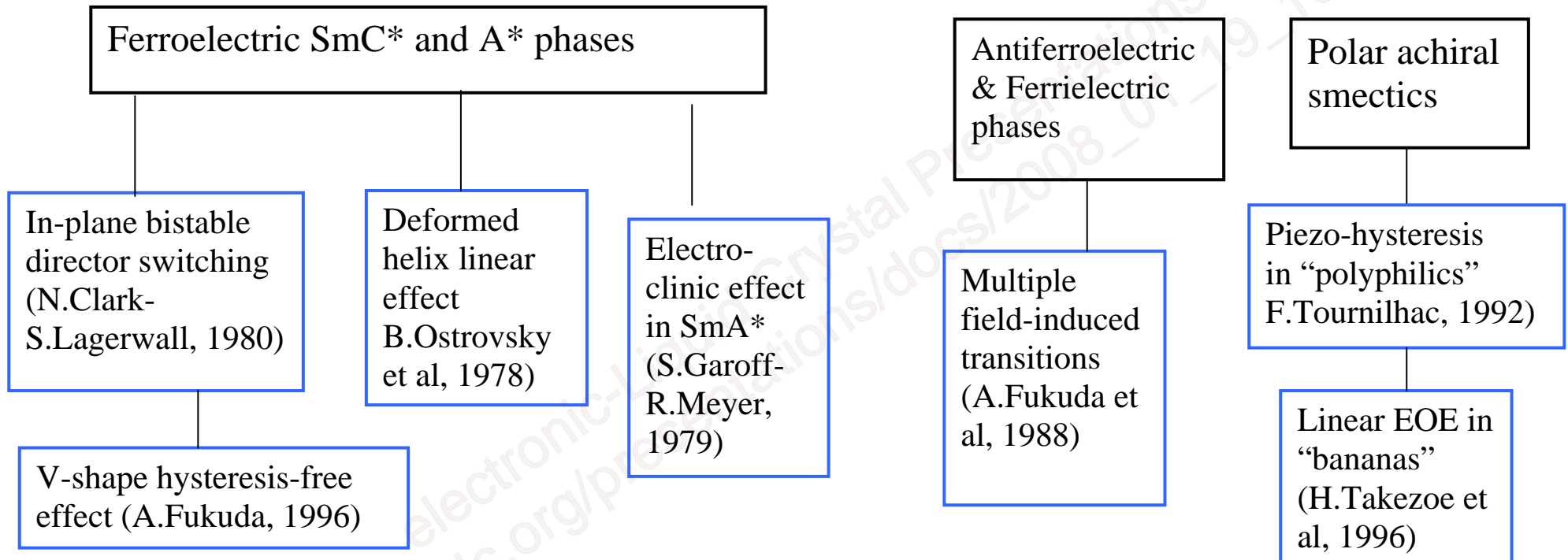


II.2 Basic EOE: Cholesterics



For applications the most important are bistable switching and weak helix distortion (dielectric and flexoelectric)

II.3 Basic EOE: Smectics



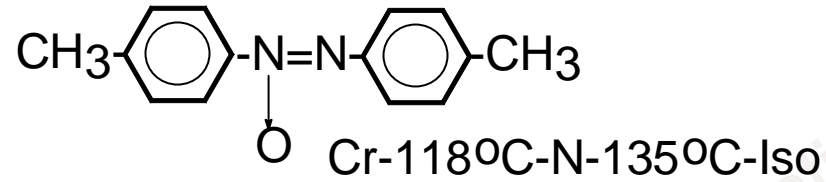
In-plane FLC switching has found application in displays and fast SLM

Two dreams:

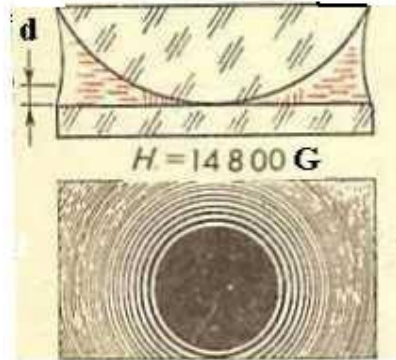
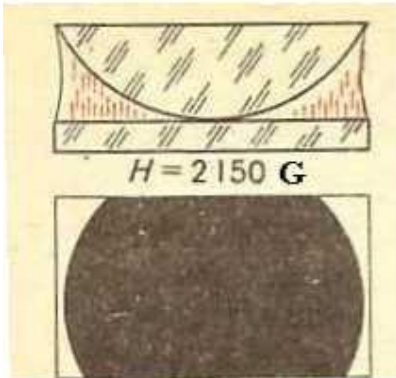
- (i) **polar nematics for linear EOE**
- (ii) **biaxial nematic for fast quadratic EOE**

III. Frederiks transition (1929)

-*p,p'*-azoxyanisole (PAA), homeotropic



V.K. Frederiks 1885-1944

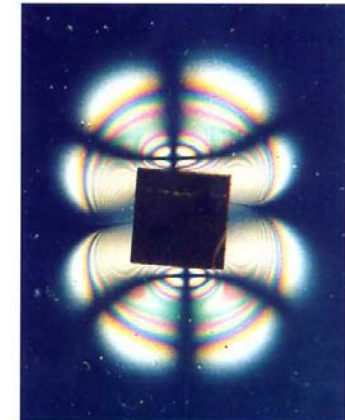
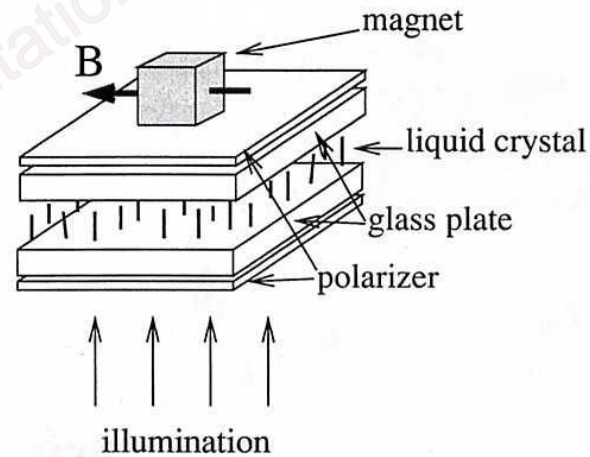


Frederiks experiment

Shown: $H_c d = \text{const.}$

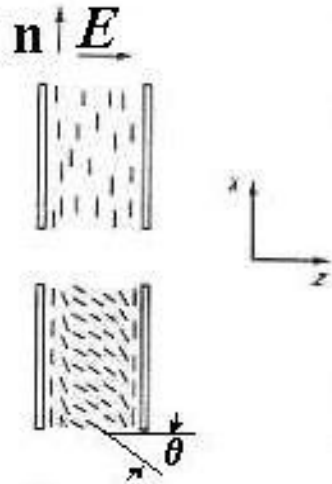
- H-field horizontal
- crossed polarizers
- Newton rings observed

Modern experiment



Simplest theory, just a balance of the elastic and electric field torques (A. Saupe (1960))

Geometry: $n_x = \cos\vartheta(z)$ and $n_z = \sin\vartheta(z)$.



Frank elastic energy:

$$g_K = \frac{1}{2} [K_{11}(\text{div}\mathbf{n})^2 + K_{22}(\mathbf{n} \cdot \text{curl}\mathbf{n})^2 + K_{33}(\mathbf{n} \times \text{curl}\mathbf{n})^2]$$

Vector algebra:

Splay: $\text{div}\mathbf{n} = \partial n_z / \partial z = \cos\vartheta \cdot (\partial\vartheta / \partial z)$

Bend: $\text{curl}\mathbf{n} = -\sin\vartheta(z) \cdot (\partial\vartheta / \partial z) \mathbf{j}$

$$\mathbf{n} \times \text{curl}\mathbf{n} = \sin\vartheta \cdot (\partial\vartheta / \partial z) \cdot (\sin\vartheta \cdot \mathbf{i} - \cos\vartheta \cdot \mathbf{k})$$

Result: $g_K = (K_{11} \cos^2 \vartheta + K_{33} \sin^2 \vartheta) \cdot \left(\frac{\partial\vartheta}{\partial z}\right)^2$ or

$$g_K = K \left(\frac{\partial\vartheta}{\partial z}\right)^2 \text{ for } K_{11} = K_{33} = K$$

Now, density of electric field energy: $g_E = \frac{\epsilon_a (\mathbf{E}\mathbf{n})^2}{4\pi}$

$$F = \frac{1}{2} \int_0^d \left[K \left(\frac{d\vartheta}{dz} \right)^2 - \frac{\varepsilon_a E^2}{4\pi} \cos^2 \vartheta \right] dz \quad \text{Minimum of } F? \text{ A variational problem:}$$

Euler equation, general: $\frac{\partial F}{\partial \vartheta} - \frac{\partial}{\partial z} \frac{\partial F}{\partial \vartheta'} = 0$ and particular: $2K \frac{\partial}{\partial z} \cdot \frac{\partial \vartheta}{\partial z} + \frac{\varepsilon_a E^2}{2\pi} \sin \vartheta \cos \vartheta = 0$

Balance of torques: $\xi^2 \frac{\partial^2 \vartheta}{\partial z^2} + \sin \vartheta \cdot \cos \vartheta = 0$ $\xi = \frac{1}{E} \sqrt{\frac{4\pi K}{\varepsilon_a}}$ **E-coherence length**

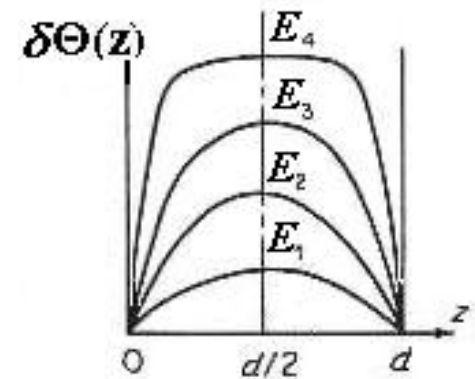
Solution in elliptic functions (A.Saupe, 1960).

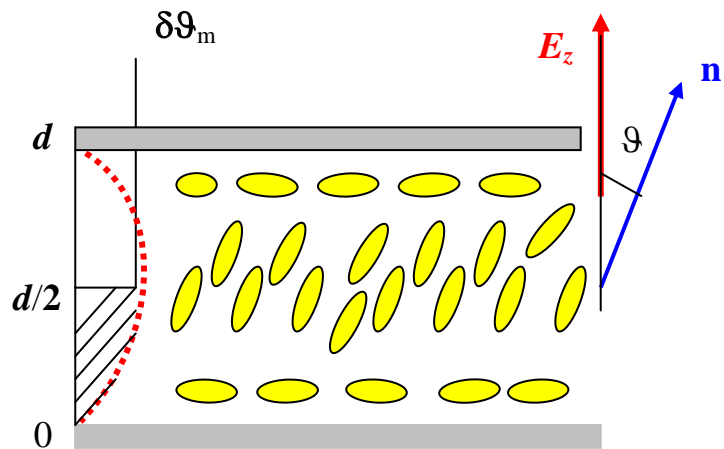
Too complicated?... Try small distortions, $\sin \vartheta \approx \vartheta$, $\cos \vartheta \approx 1$

$$\xi^2 \frac{\partial^2 \vartheta}{\partial z^2} + \vartheta = 0 \quad \text{linear equation with harmonic functions}$$

$$\delta \vartheta = \pi/2 - \vartheta = \delta \vartheta_m \sin qz \quad \text{with } q = \pi/d.$$

After substitution: $-\xi^2 q^2 + 1 = 0$ or $\xi_{Fr} = \frac{d}{\pi}$





Threshold condition: $\pi \xi_{Fr} = d$ or $E_{Fr} = \frac{\pi}{d} \sqrt{\frac{4\pi K}{\epsilon_a}}$

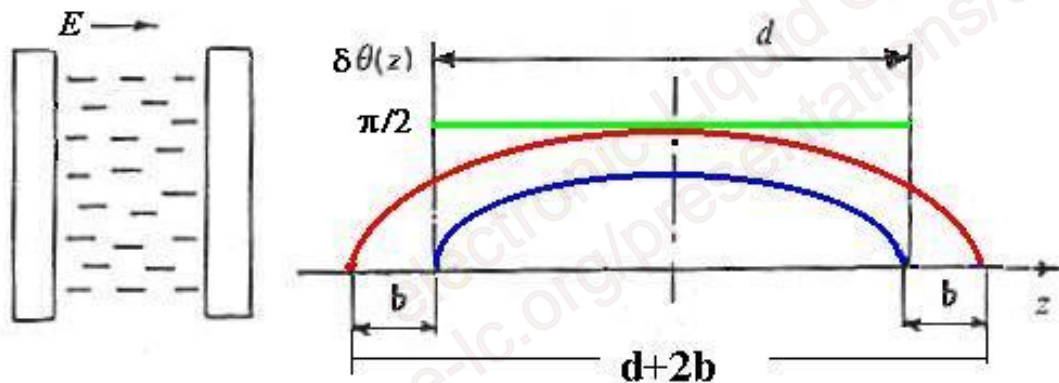
Frederiks experiment: $H_c d = \text{const}$

Weak anchoring energy W^s

and new characteristic length $b = K_{ii}/W^s$

Break of anchoring- new threshold:

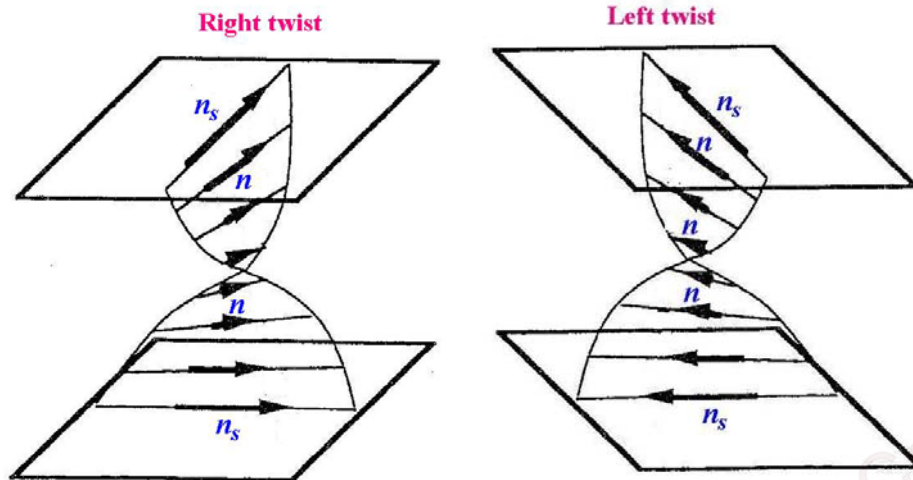
$$\pi \xi_{BA} = b$$



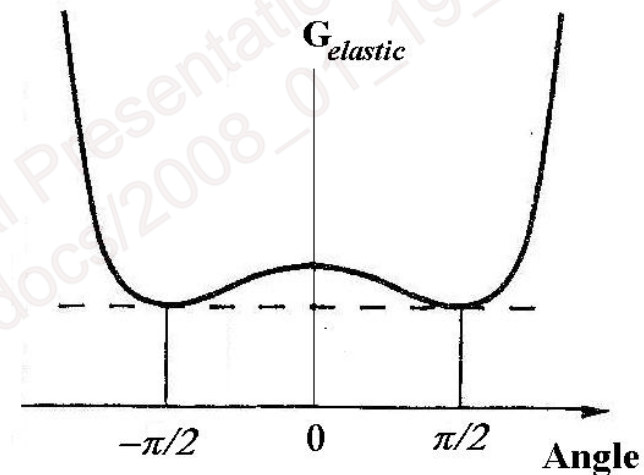
relax? Frustration: the idea for bistable switching!

After the break: which way to

The naive idea of bistability for a twisted nematic:



For strongly rubbed plates the twisted structure has lower energy $F_{el} + F_s$

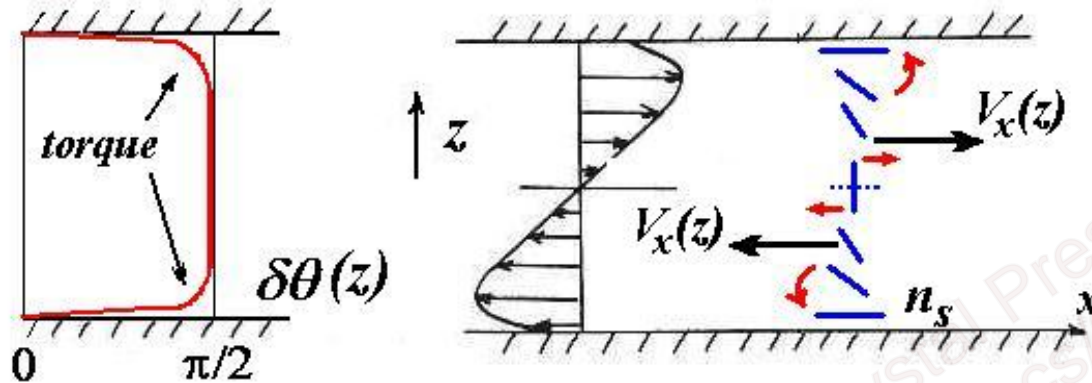


The two structures transmit CPL differently.

But how to force the director select one of them by field application?

First to orient director almost vertically and then to let it relax to one of the two structures using a torque of non-electric nature, e.g. hydrodynamic (D.Berreman, G.Durand, R.Barberi, F.Martinot-Lagarde, I.Dofov, S.Palto)...

Backflow effect:



Important:

Fast relaxation of the director close to the plates creates a flow, which, in turn, drives the director in the middle of the cell

This can be used for the final

structure selection

$$\rho \frac{\partial V}{\partial t} = \frac{\partial}{\partial z} \left(A \frac{\partial \vartheta}{\partial t} + B \frac{\partial V}{\partial z} \right)$$

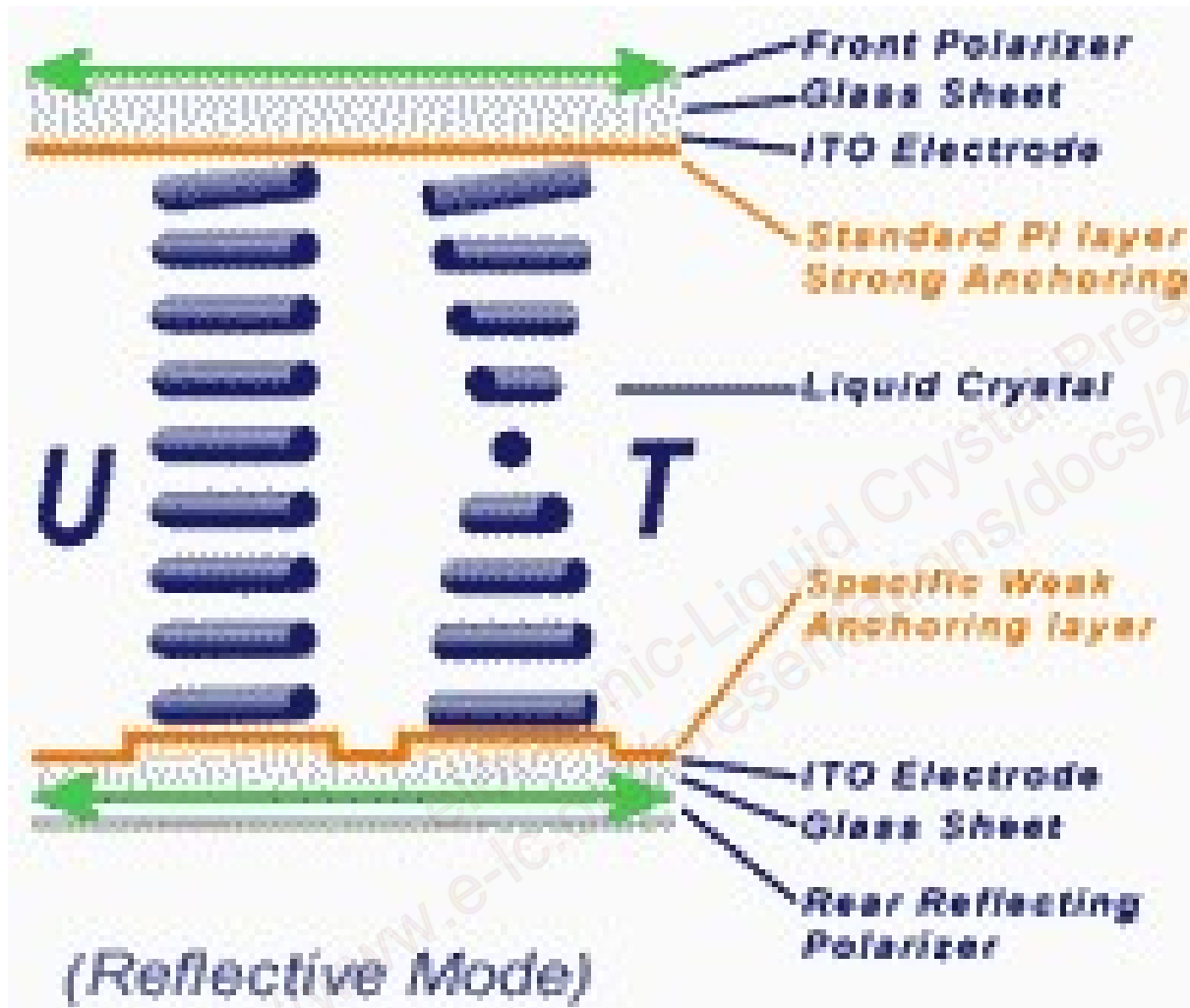
equation of motion for liquid

A, B – combinations of viscosities

$$\gamma_1 \frac{\partial \vartheta}{\partial t} = K \frac{\partial^2 \vartheta}{\partial z^2} + \frac{\varepsilon_a}{4\pi} \sin \vartheta \cos \vartheta - A \frac{\partial V}{\partial z}$$

torque balance for the director

Bistable Nematic display with memory

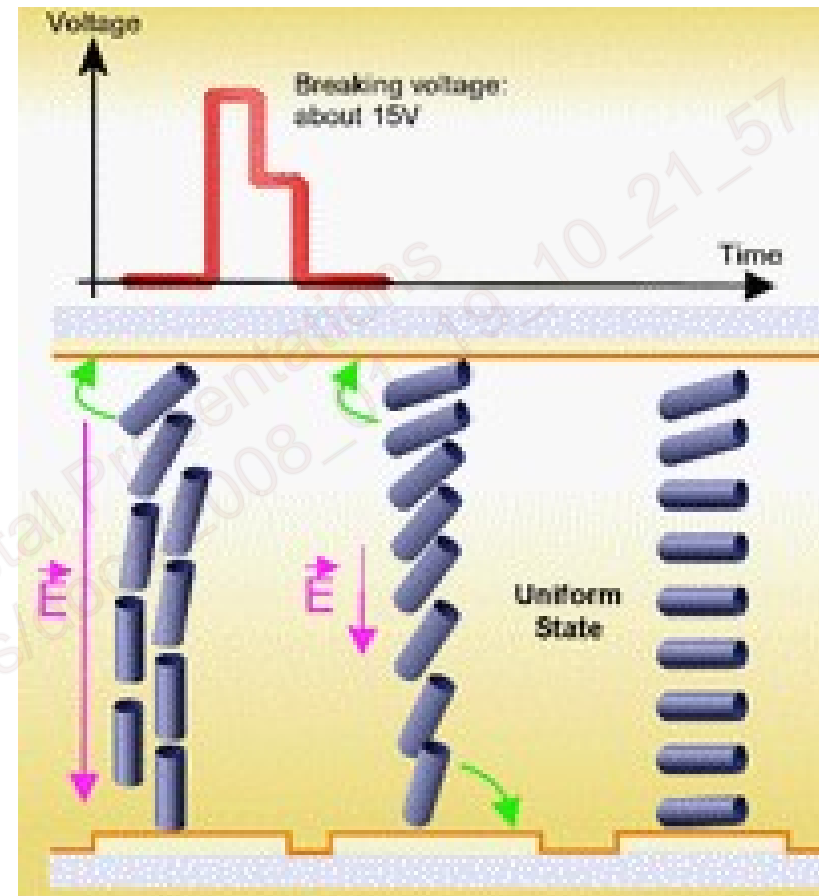
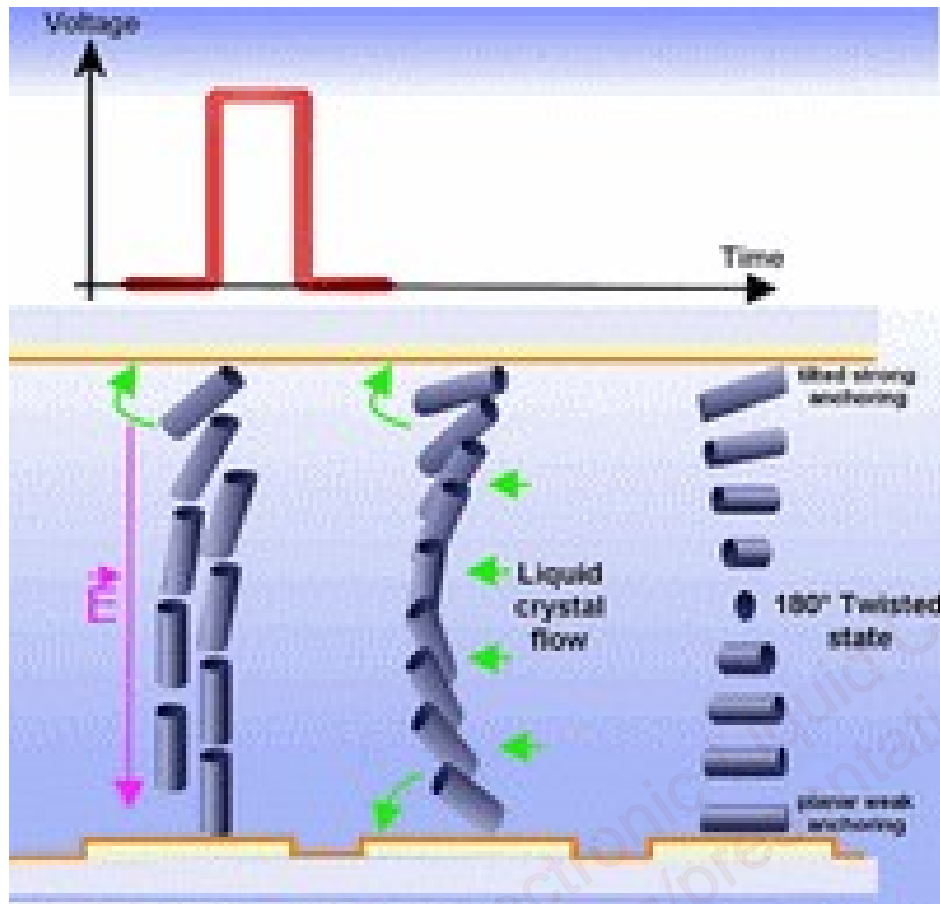


LC has well tuned (weak) chirality

The two structures have similar minima of elastic energy

The director position is different in the cell center

The backflow selects the T-structure during relaxation after the break of anchoring.



Steep rear edge of pulse causes strong twisting backflow

Step-like edge results in smooth relaxation to uniform state

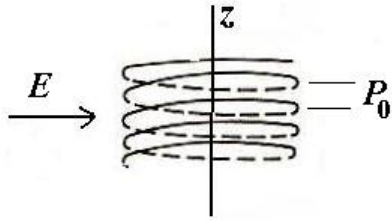
The result is a beautiful display with memory for energy saving devices, smart cards, electronic paper, electronic posters for the next OLC conferences, etc)



http..

presentations
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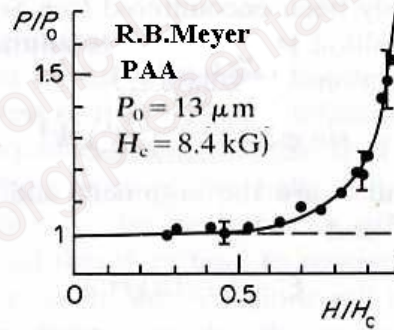
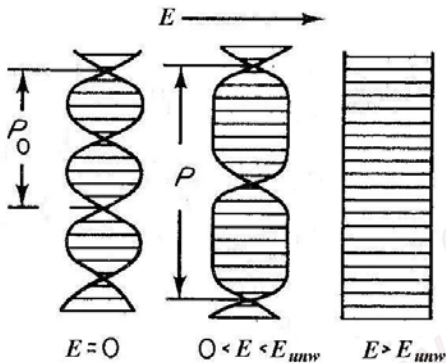
IV. Helix unwinding (P.G. de Gennes, R.B. Meyer, 1968)



$$g_{Ch} = \frac{1}{2} \left[K_{22} \left(\frac{d\varphi}{dz} - q_0 \right)^2 - \frac{\epsilon_a E^2}{4\pi} \cos^2 \varphi \right] \quad (\text{infinite helix})$$

Unwinding threshold: $E_{unw} = \frac{\pi^2}{P_0} \sqrt{\frac{4\pi K_{22}}{\epsilon_a}}$ or $\pi^2 \xi_{unw} = \frac{2\pi}{q_0} = P_0$

Field dependence of pitch: $P(E) = P_0 \left[1 + \frac{\epsilon_a^2 P_0^2}{2^{11} \pi^5 K_{22}^2} E^4 + \dots \right]$

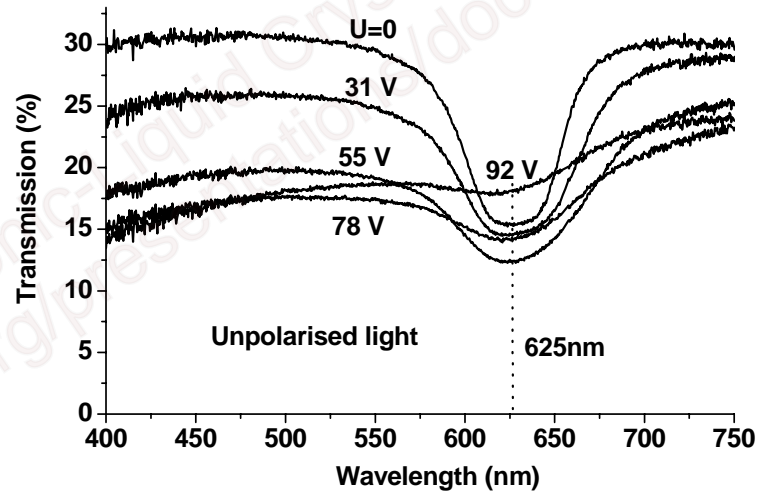
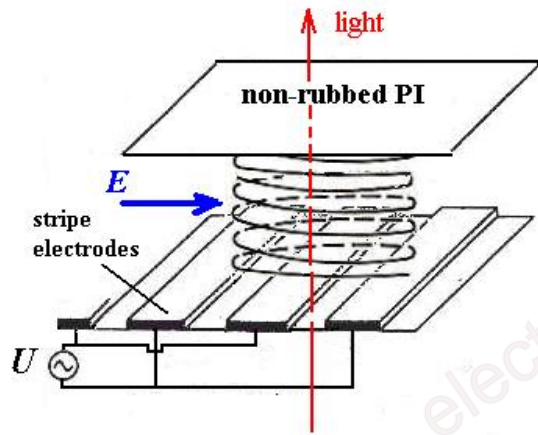
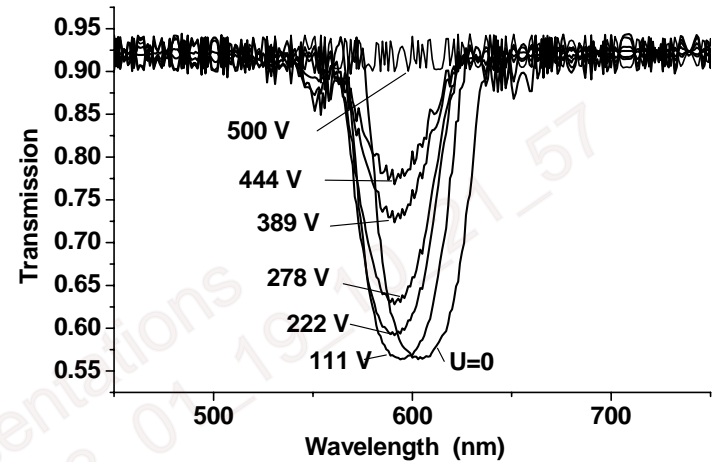
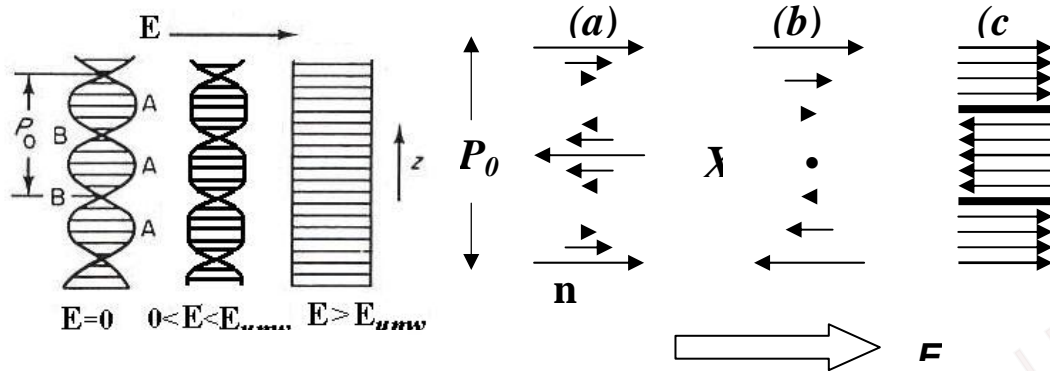


Hopes

Fast effect for short pitch $\tau = \frac{\gamma_1}{K_{22} q^2} \quad q=2\pi/P$

- fast color displays
- field controllable optical diffraction,
- tunable DFB lasers, etc

More realistic situation: modelling and experiment (S. Palto, L.Blinov 2005)

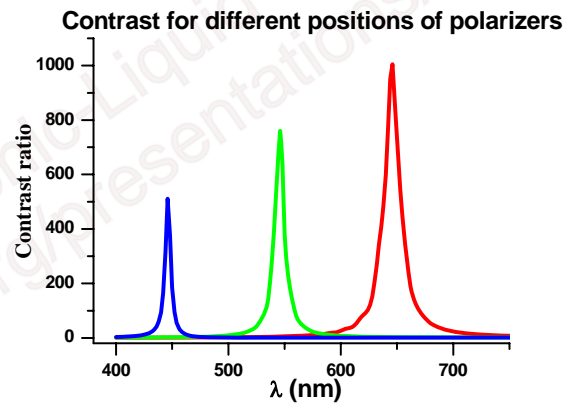
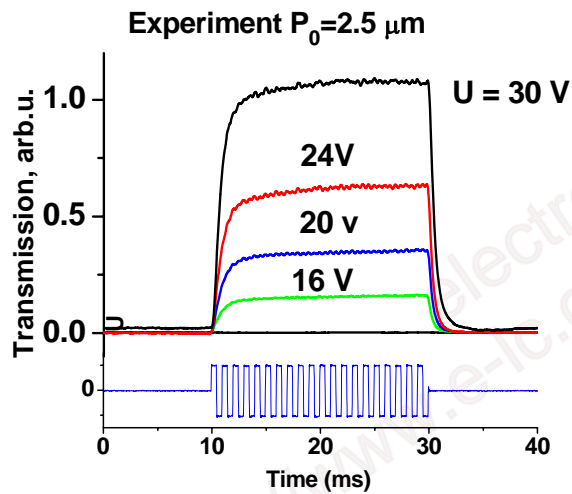
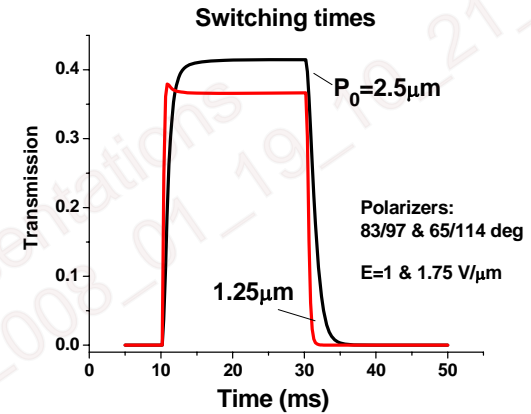
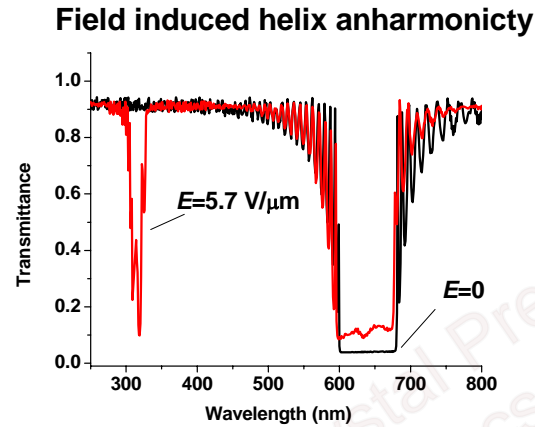
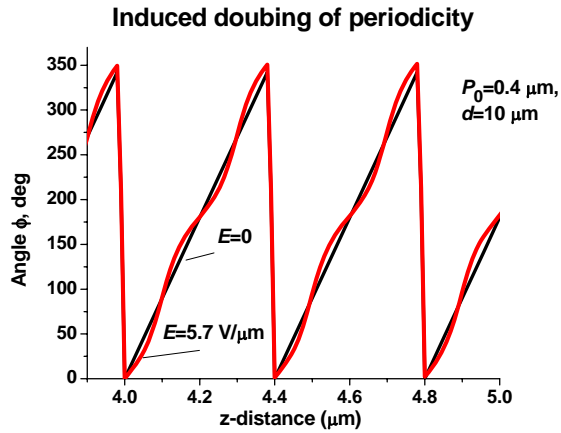


Thermodynamics is not sufficient, there are topological confinements even for infinite helix

The helix can be unwound and wound back only very slowly, via defects

Usually one observe only field induced anharmonicity without change of the period

Field induced anharmonicity and fast display (M.Barnik, S.Palto et al, 2006)



Results

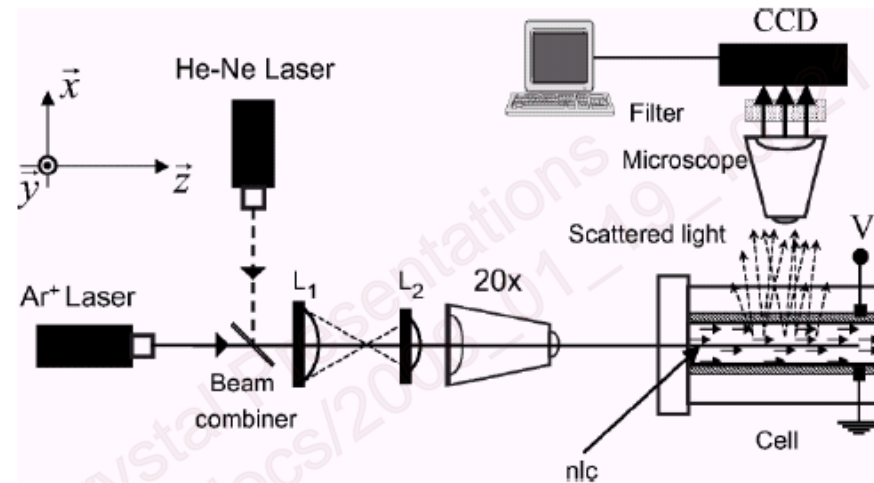
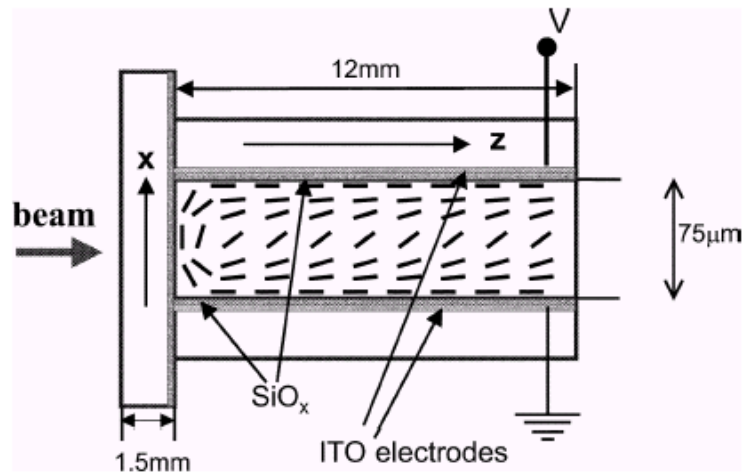
Times: 4 times shorter

(400 μs -1.5 ms) due to doubling of period of the structure:

$$\tau = \frac{\gamma_1}{4K_{22}q^2}$$

High contrast is solely defined by polarizers

V. Hybrid of electro-optics with non-linear optics: field control of spatial solitons

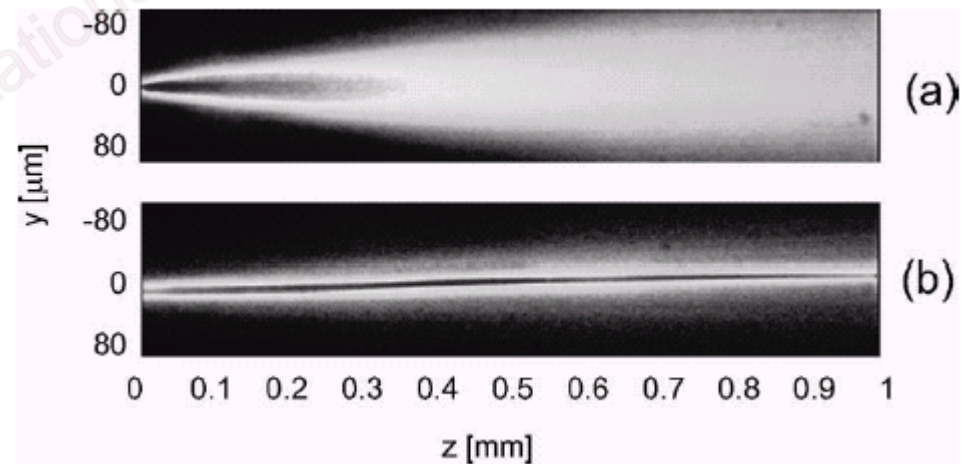


XXIII. G. Assanto et al

Idea: To use the non-linear effect of beam self-focusing for compensation of the beam divergence.

Experiment:

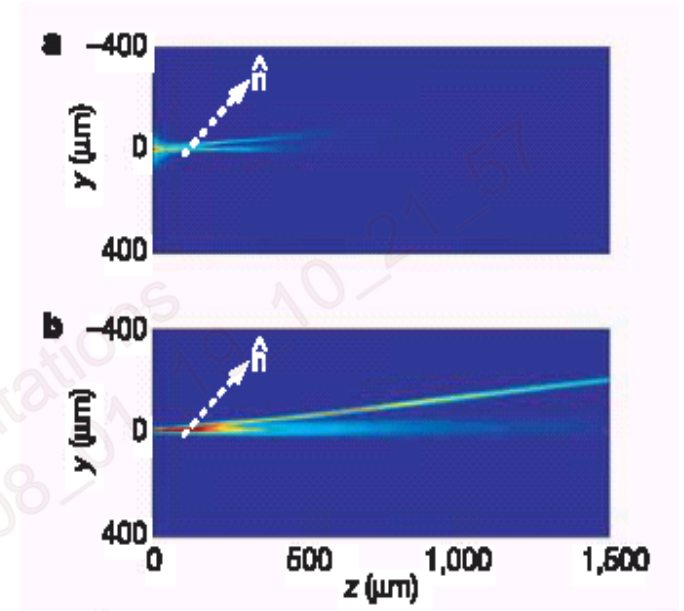
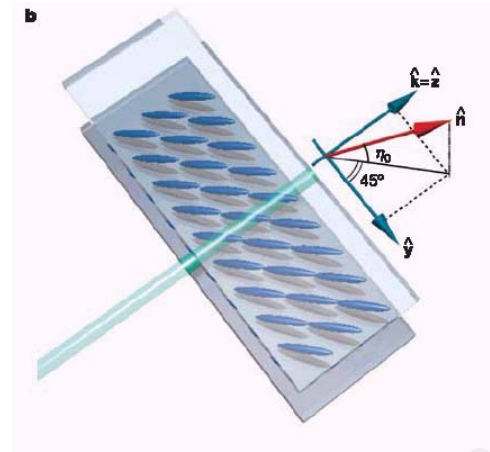
$U=1\text{V}$, y - (a) and x - (b) polarized beams
IEEE J. Quant. Electr. 39, 13 (2003)



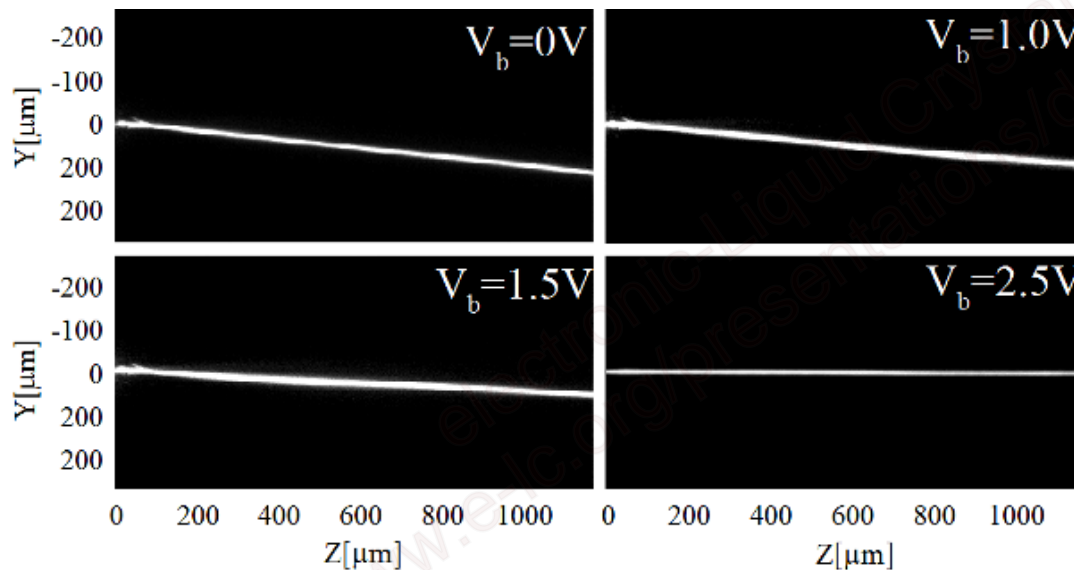
Routing of anisotropic spatial solitons and modulational instability in liquid crystals

Marco Peccianti¹, Claudio Conti¹, Gaetano Assanto¹, Antonio De Luca² & Cesare Umeton²

Nature, 432, 734 (2004)



Zero bias and low beam power (a); At higher beam power (b) e-beam generates self-waveguiding soliton



The direction of soliton propagation is controlled by voltage

Take home:

Electrooptics of liquid crystals is mature science with hundreds of effects discovered and understood

Only few effects are used in conventional display technology, mostly based on the different versions of the Frederiks transition

There are serious demands for new effects for future display technology (smart cards, electronic paper, etc.), optics (fast SLM, field controlled diffraction elements, waveguiding devices) and nonlinear optics (laser devices, phase conjugation, routing, etc.).

Acknowledgements: to all my colleagues from the Calabria University and the Institute of Crystallography of Russian Academy of Sciences